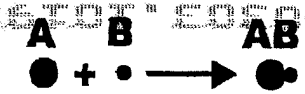
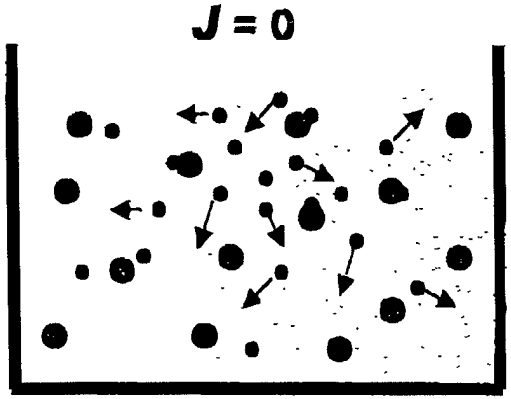
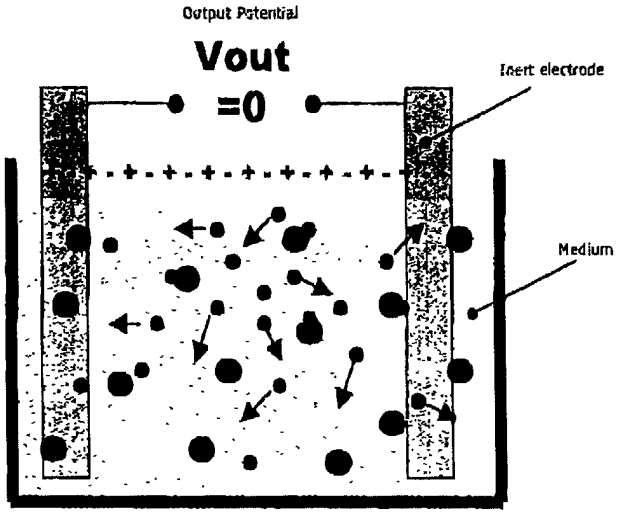


J : Current Density of B

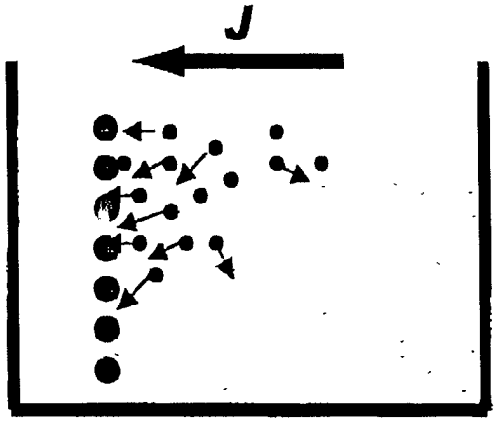
1.1

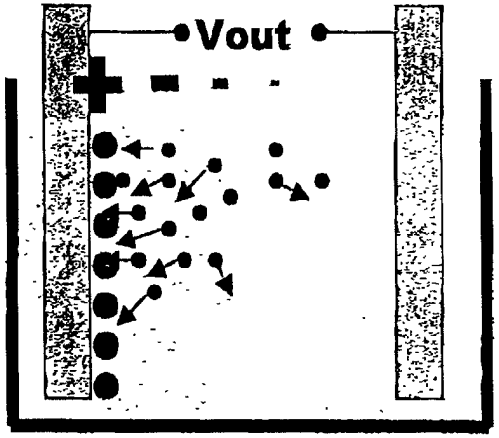




1.1) If A and B are both free in the medium no net transient gradient of molecules (current density) is created.

1.2

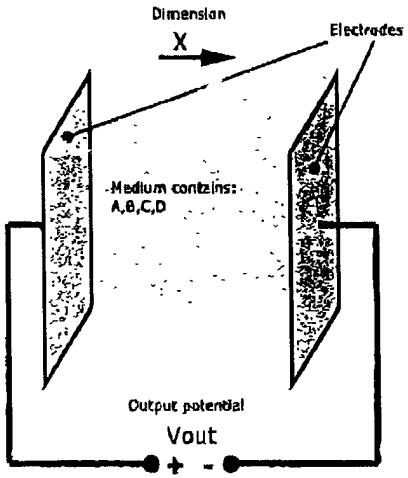


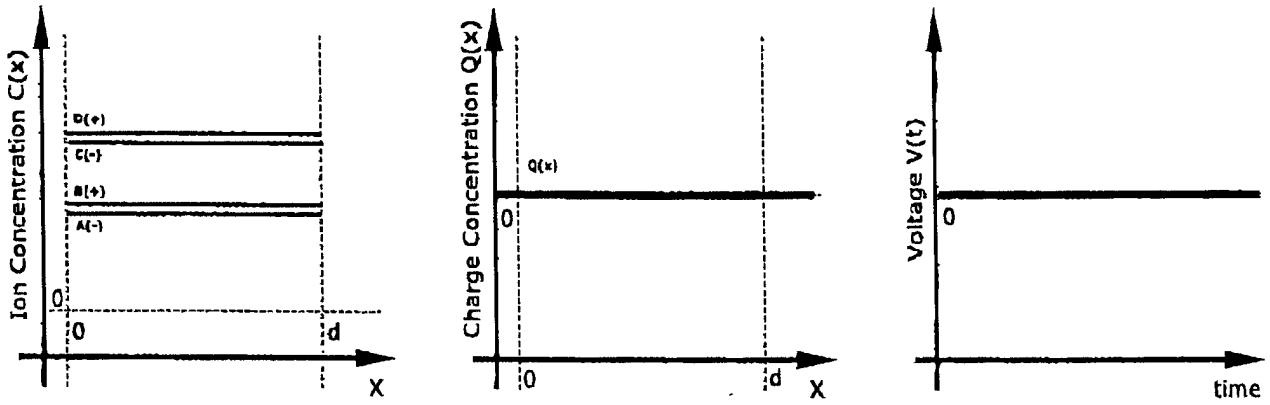


1.2) If A is spatially immobilized and B is free in the medium, the reaction causes a net transient gradient (current density) of B toward A. This transient current creates a temporary potential difference in the medium.

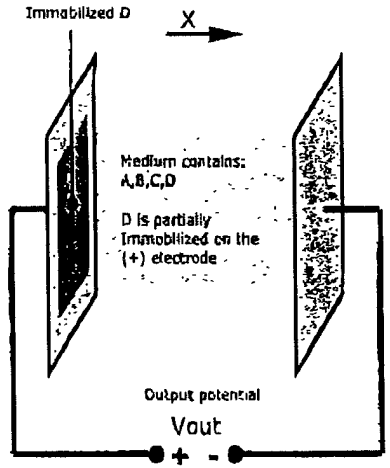
1.3 Potential difference between electrodes equilibrium (steady state):

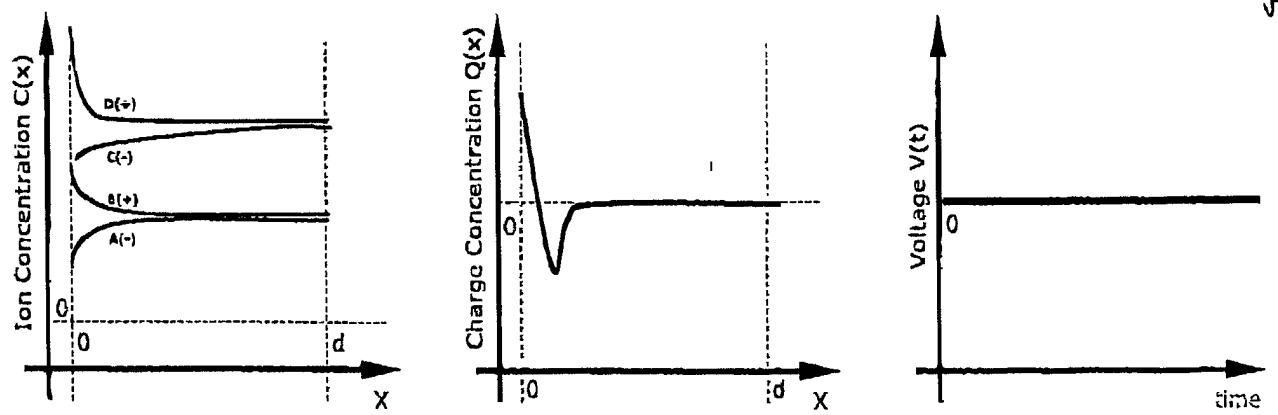
1) Electrodes are inert and do not interact with medium.
2) Four molecules (ions) A, B, C and D are present in the medium.
3) Molecules have arbitrary diffusion length and charge.





1.3.a) All molecules are free in the medium. Concentration of all molecules is uniform in equilibrium.

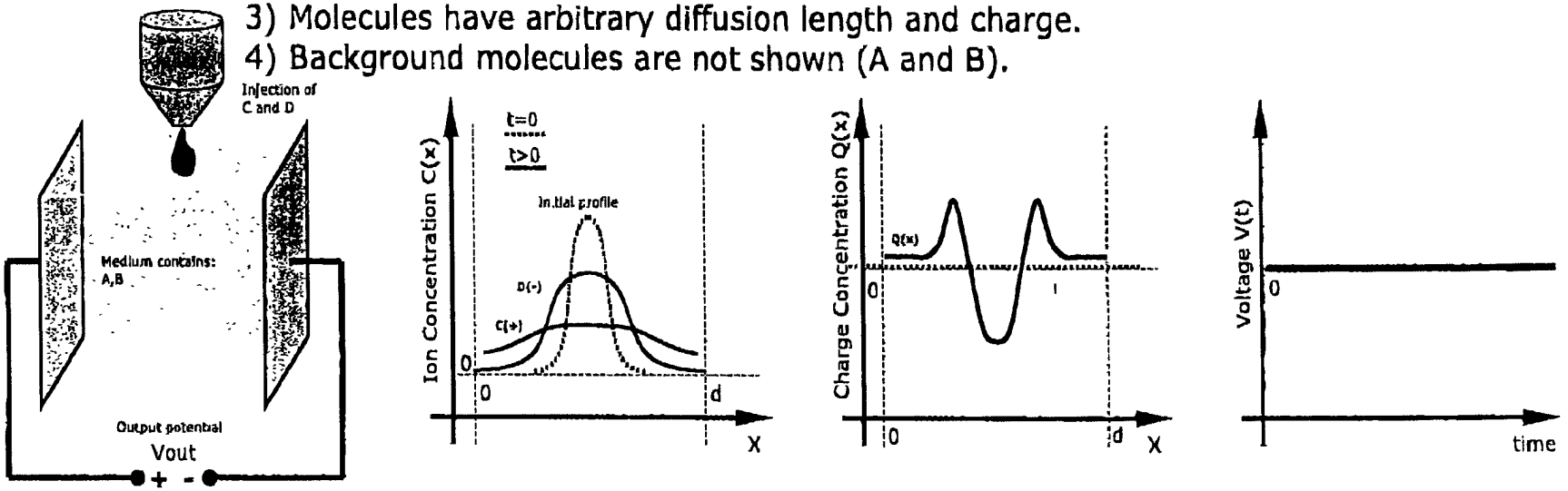




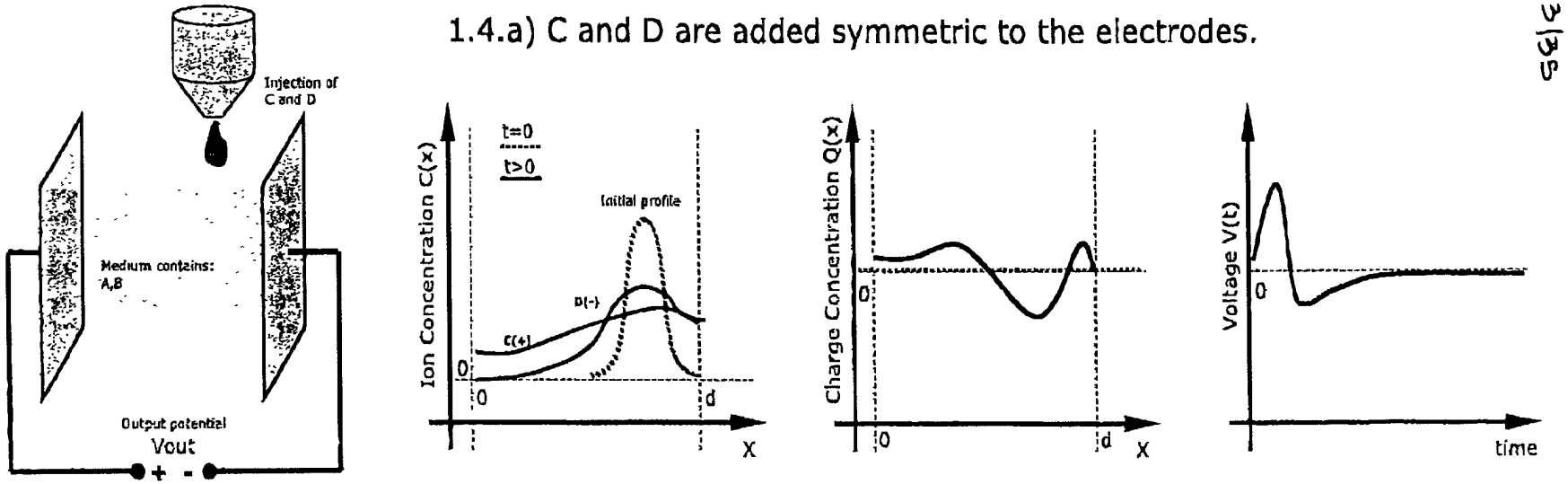
1.3.b) D is partially immobilized on the surface of the (+) Electrode which forces a none uniform concentration of molecules.

1.4 Potential difference between electrodes in none-equilibrium (transient) state:

1) Electrodes are inert and do not interact with the medium.
2) Two molecules (ions) are present in the medium and two are added.
3) Molecules have arbitrary diffusion length and charge.
4) Background molecules are not shown (A and B).



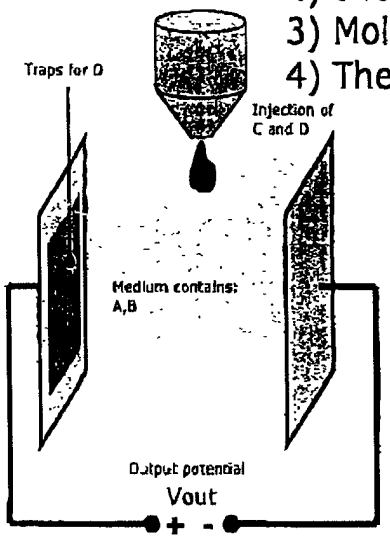
1.4.a) C and D are added symmetric to the electrodes.

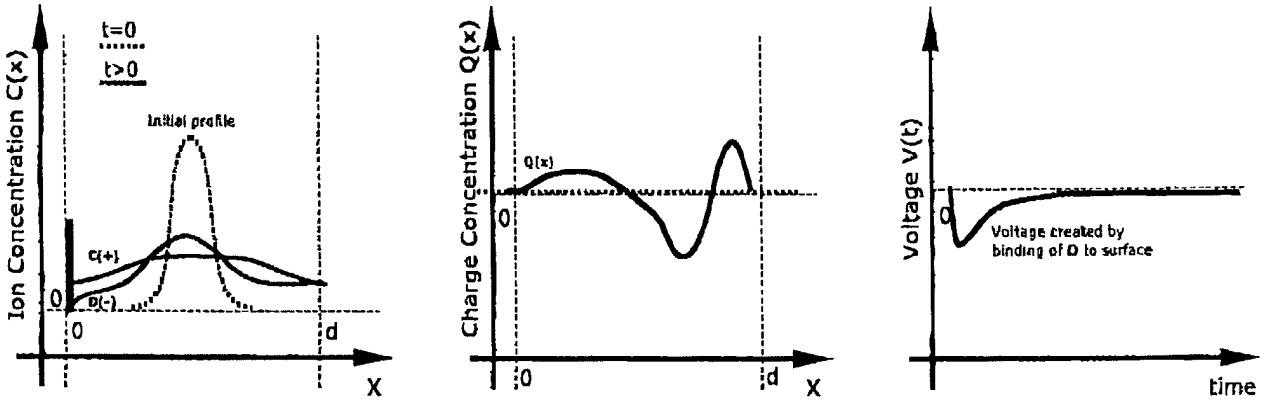


1.4.b) C and D are added asymmetric to the electrodes and an ionic perturbation is generated.

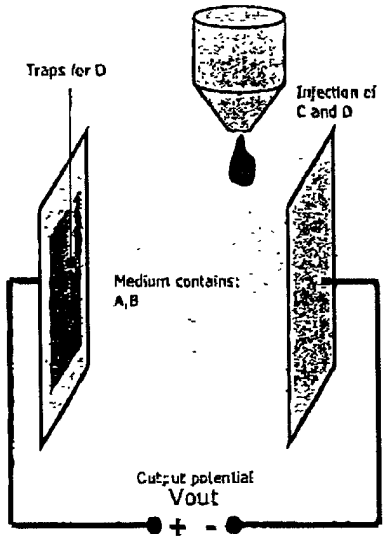
1.5 Potential difference between electrodes in the none-equilibrium (transient) with surface trap:

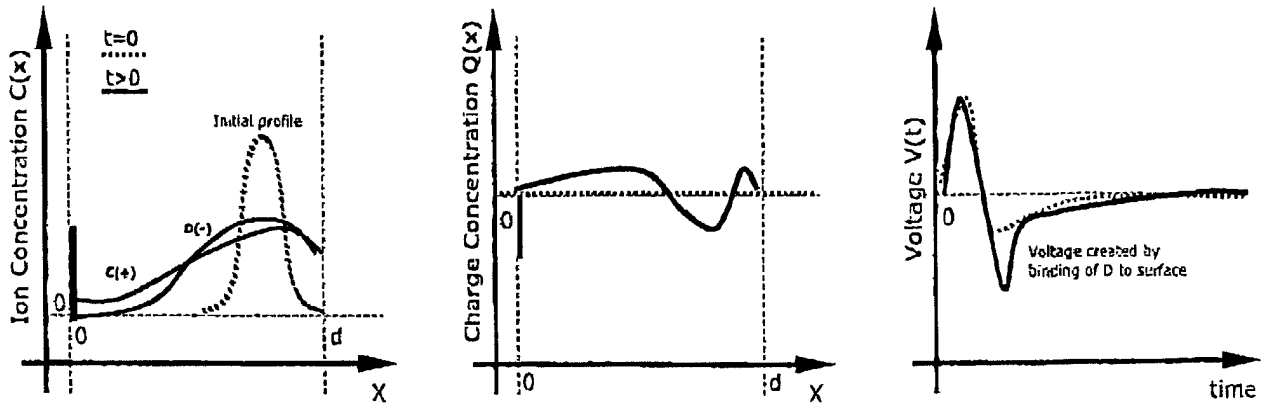
1) Electrodes are inert and do not interact with the medium.
2) two molecules (ions) are present in the medium and two are added in time.
3) Molecules have arbitrary diffusion length and charge.
4) The (+) electrode has finite traps for D on the surface.





1.5.a) C and D are added symmetric to the electrodes. Traps cause a potential perturbation.

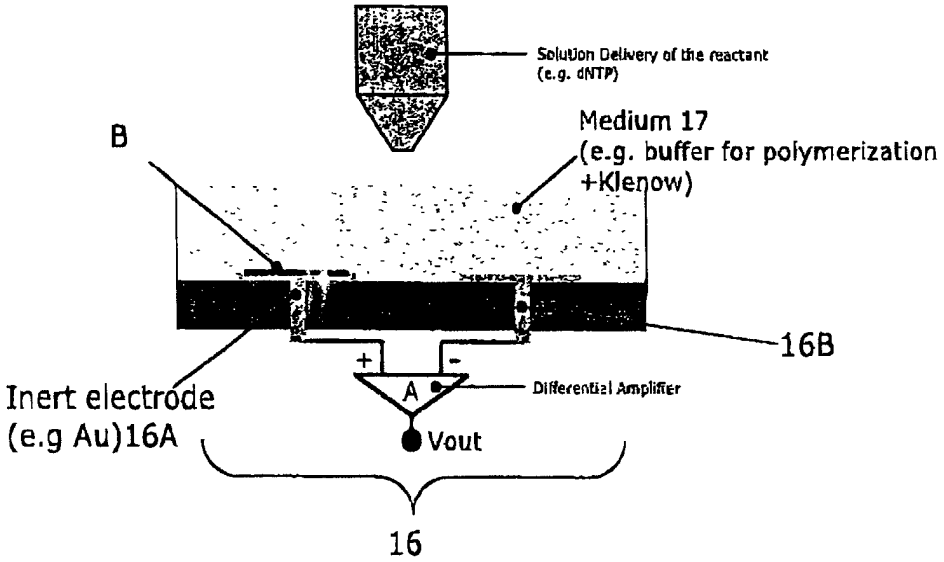




1.5.b) C and D are added asymmetric to the electrodes, and an extra electric field perturbation is created by the traps.

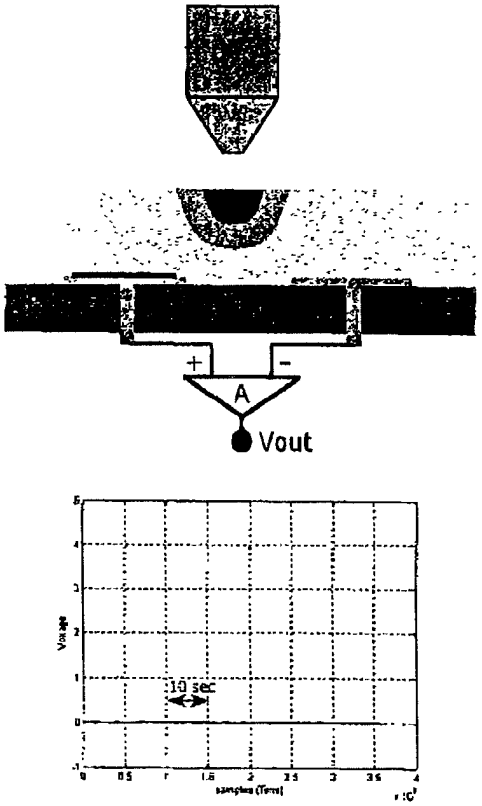
2.1 Planar sensor design example:

1) Electrodes are inert and do not interact with the medium.
2) The target molecules are immobilized on the (+) electrode.
3) The (-) electrode is the reference electrode.
4) A differential amplifier subtracts the voltage from the two electrodes.

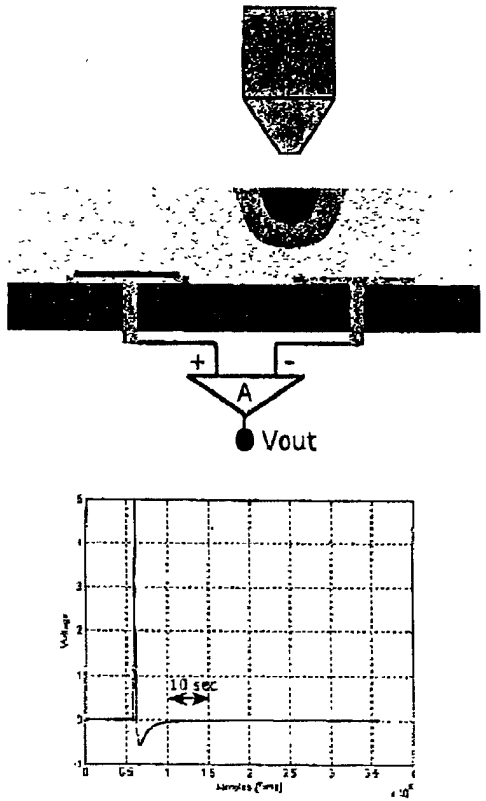


2.2 Example of signal generated when no binding at the surface occurs :

1) Electrodes are inert and do not interact with the medium.
2) The target molecules are immobilized on the (+) electrode.
3) The (-) electrode is the reference electrode.
4) A differential amplifier subtracts the voltage of the two electrodes.



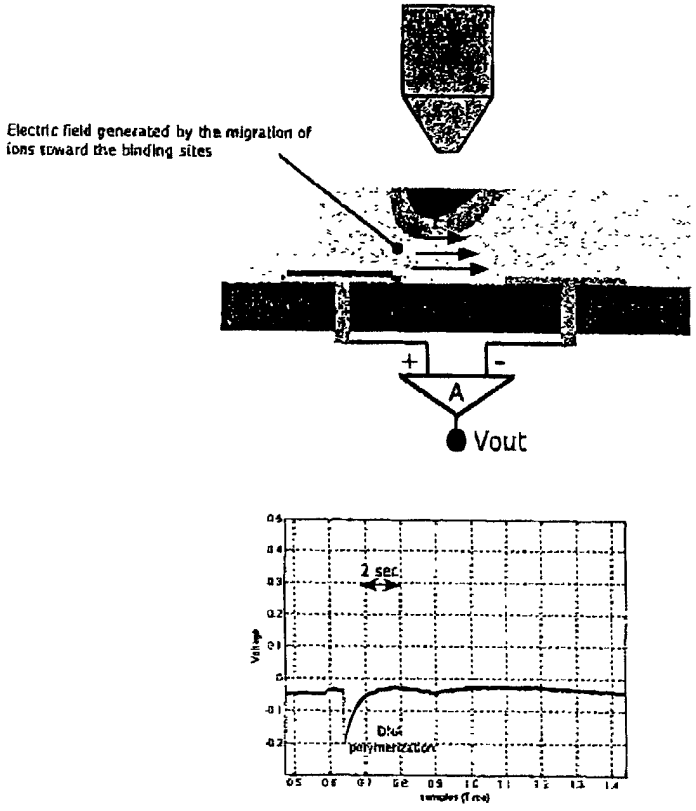
a) Solution is delivered symmetric to the electrodes With 0.1 pmol immobilized ssDNA.



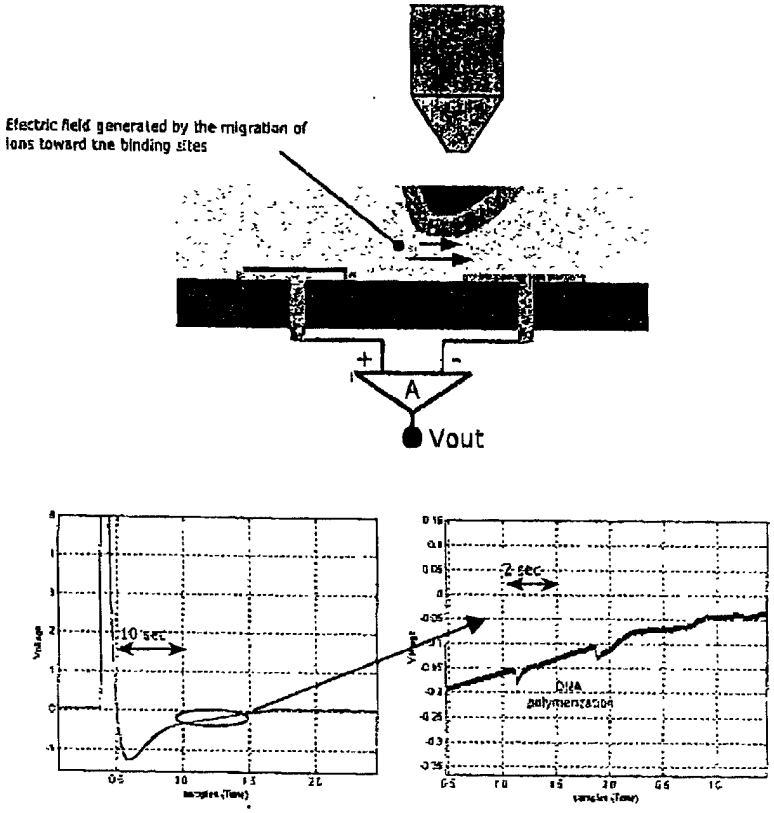
b) Solution is delivered asymmetric to the electrodes With 0.1 pmol immobilized ssDNA.

2.3 Example of signal generated when binding at the surface occurs:

1) Electrodes are inert and do not interact with the medium.
2) The target molecules are immobilized on the (+) electrode.
3) The (-) electrode is the reference electrode.
4) A differential amplifier subtracts the voltage from the two electrodes.

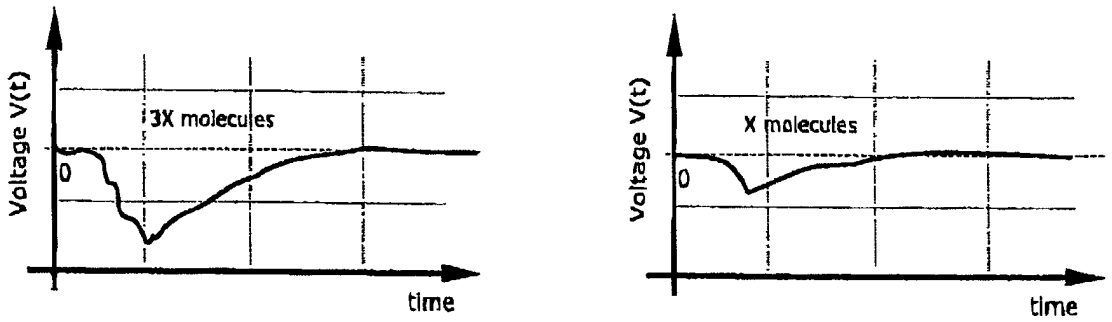


a) Solution is delivered symmetric to the electrodes, Polymerization of 0.1 pmol primed ssDNA.

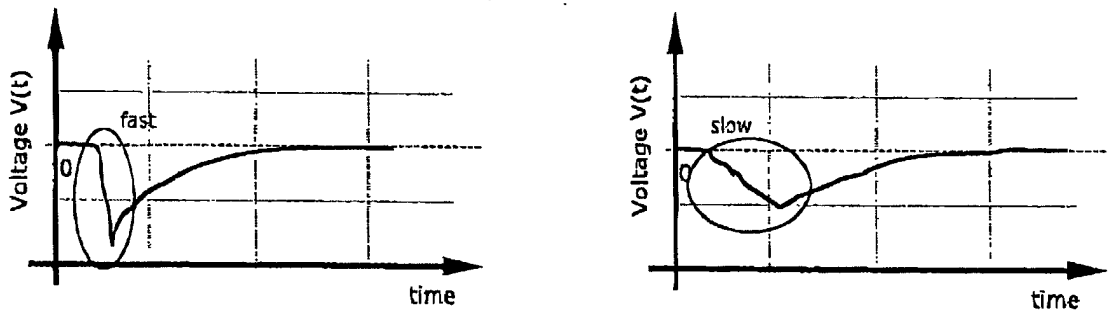


b) Solution is delivered asymmetric to the electrodes Polymerization of 0.1 pmol primed ssDNA.

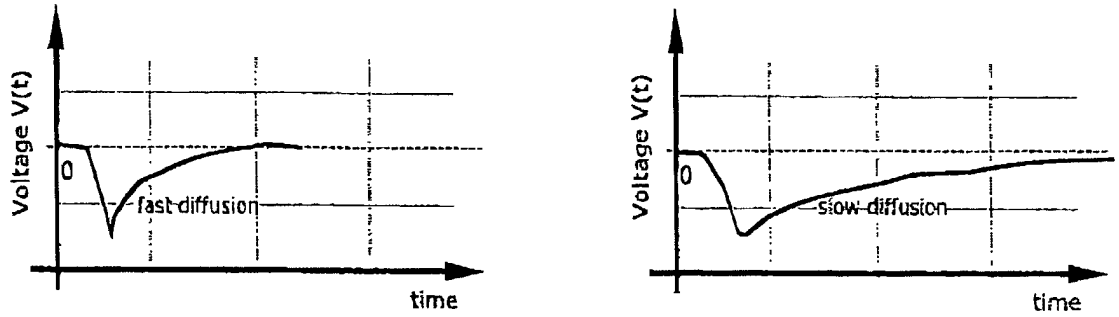
2.4 Analysis examples of the transient signal generated:



a) Quantity of molecules

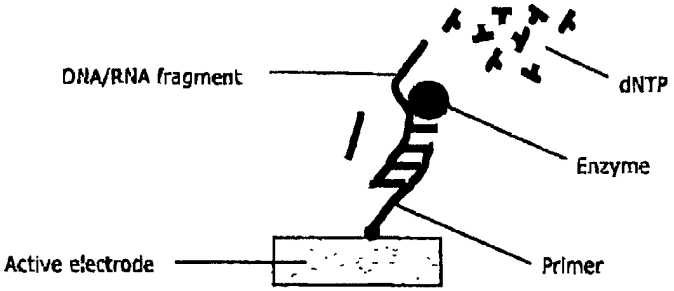


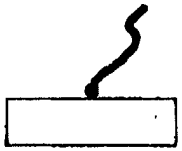
b) Kinetics (speed) of the reaction



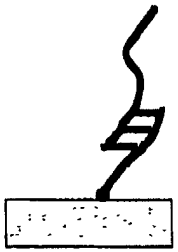
a) Movement and diffusion of molecules.

FIGURE 3:
Sequencing

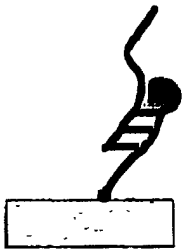




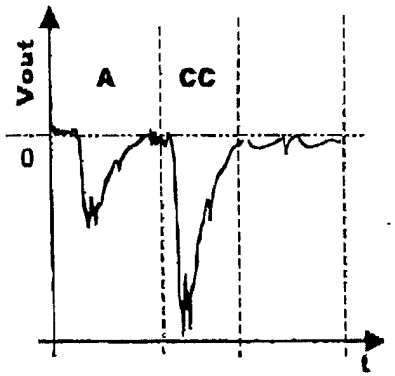
Gene specific primer

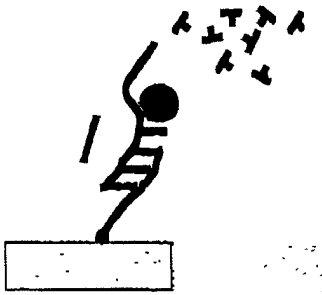


Hybridization of DNA/RNA to primer

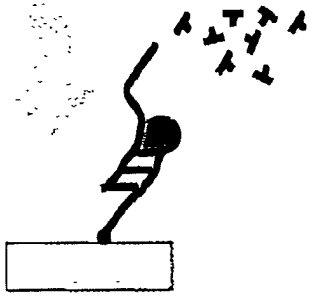


Addition of Enzyme





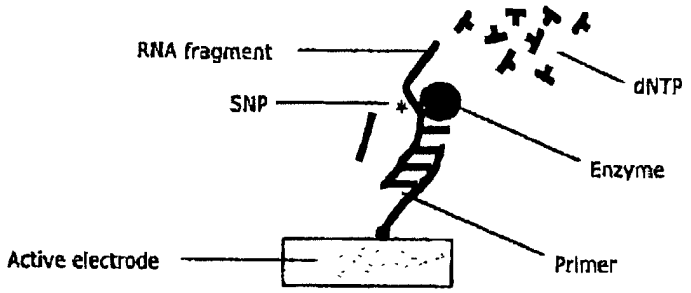
Right nucleotide added and signal generation

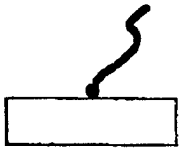


Addition of specific Nucleotide one at a time or all at the same time

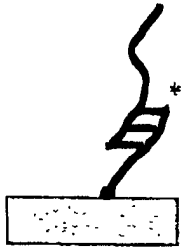
FIGURE 4:1

SNP detection by using Total RNA

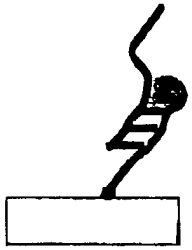




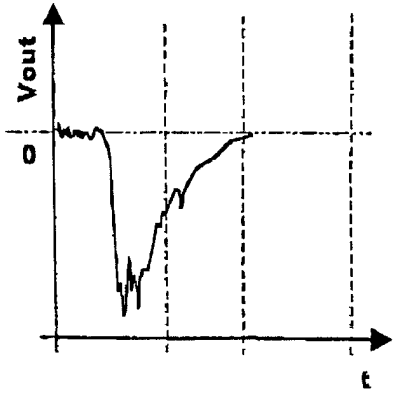
Gene specific designed primer around the SNP

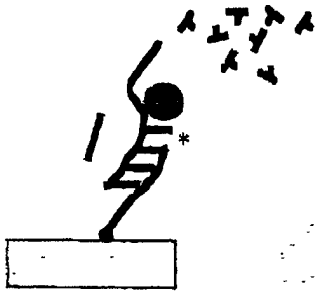


Hybridization of RNA to Specific primer

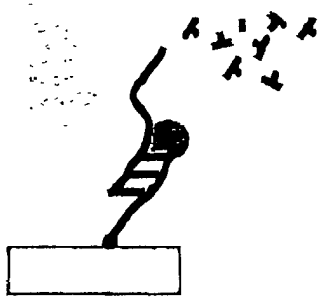


Addition of Enzyme





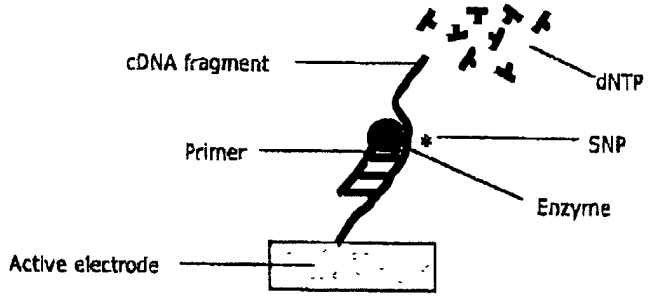
Correct nucleotide generates signal

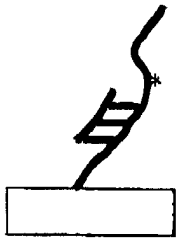


Addition of specific nucleotide

FIGURE 4:2

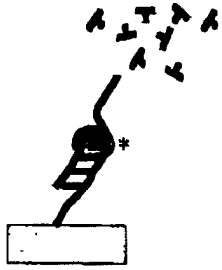
SNP detection by using cDNA



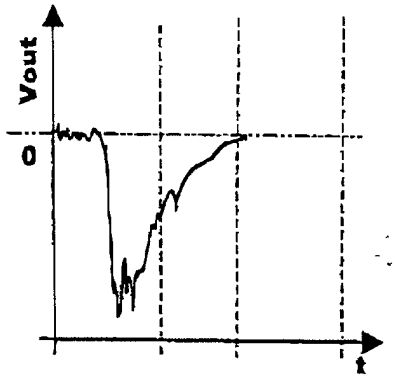


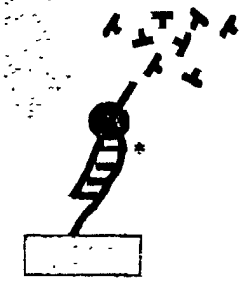
Annealing the specific designed primer around the SNP

Addition of Enzyme



Addition of specific nucleotide



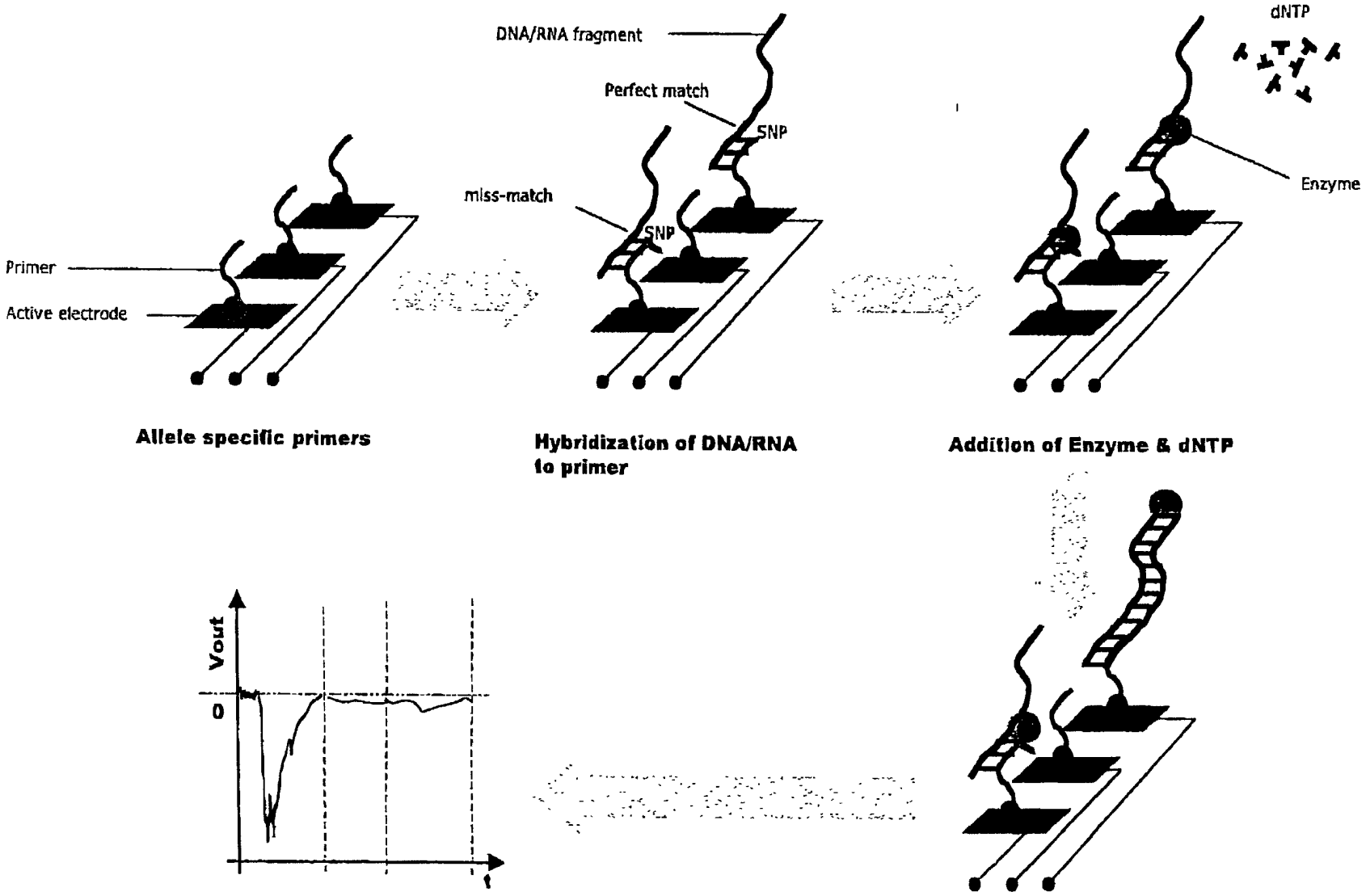


Right nucleotide added and signal generation

1135

FIGURE 4:3

SNP detection by using allele specific primer



Allele specific primers

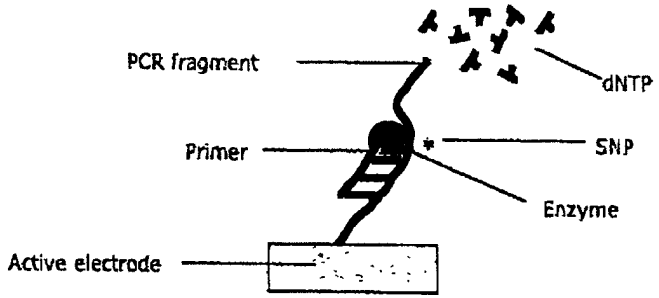
Hybridization of DNA/RNA to primer

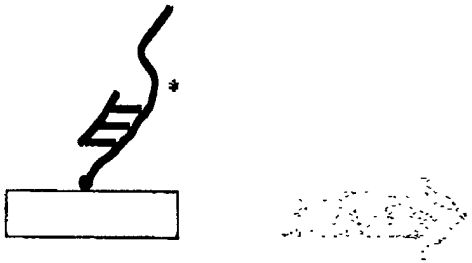
Addition of Enzyme & dNTP

Perfect match primer/template polymerizes and signal is generated

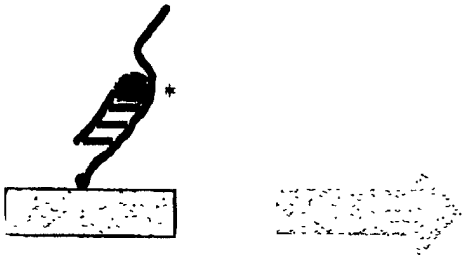
FIGURE 5:

SNP detection by using PCR product

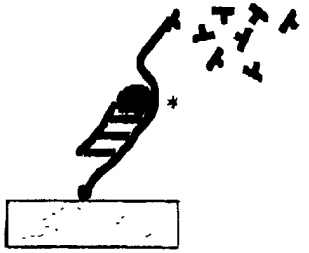




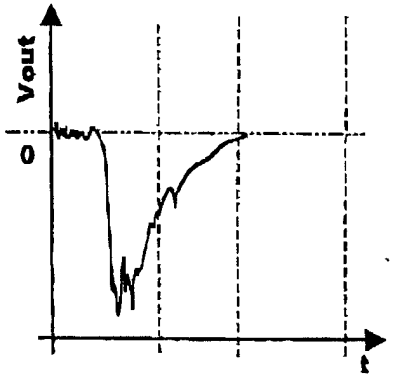
Annealing the specific designed primer around the SNP

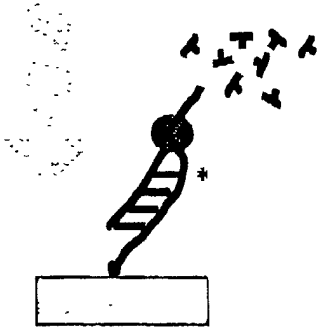


Addition of Enzyme



Addition of specific nucleotide





Right nucleotide added and signal generation

FIGURE 6:

SNP detection by using Exonuclease/ Degrading enzyme

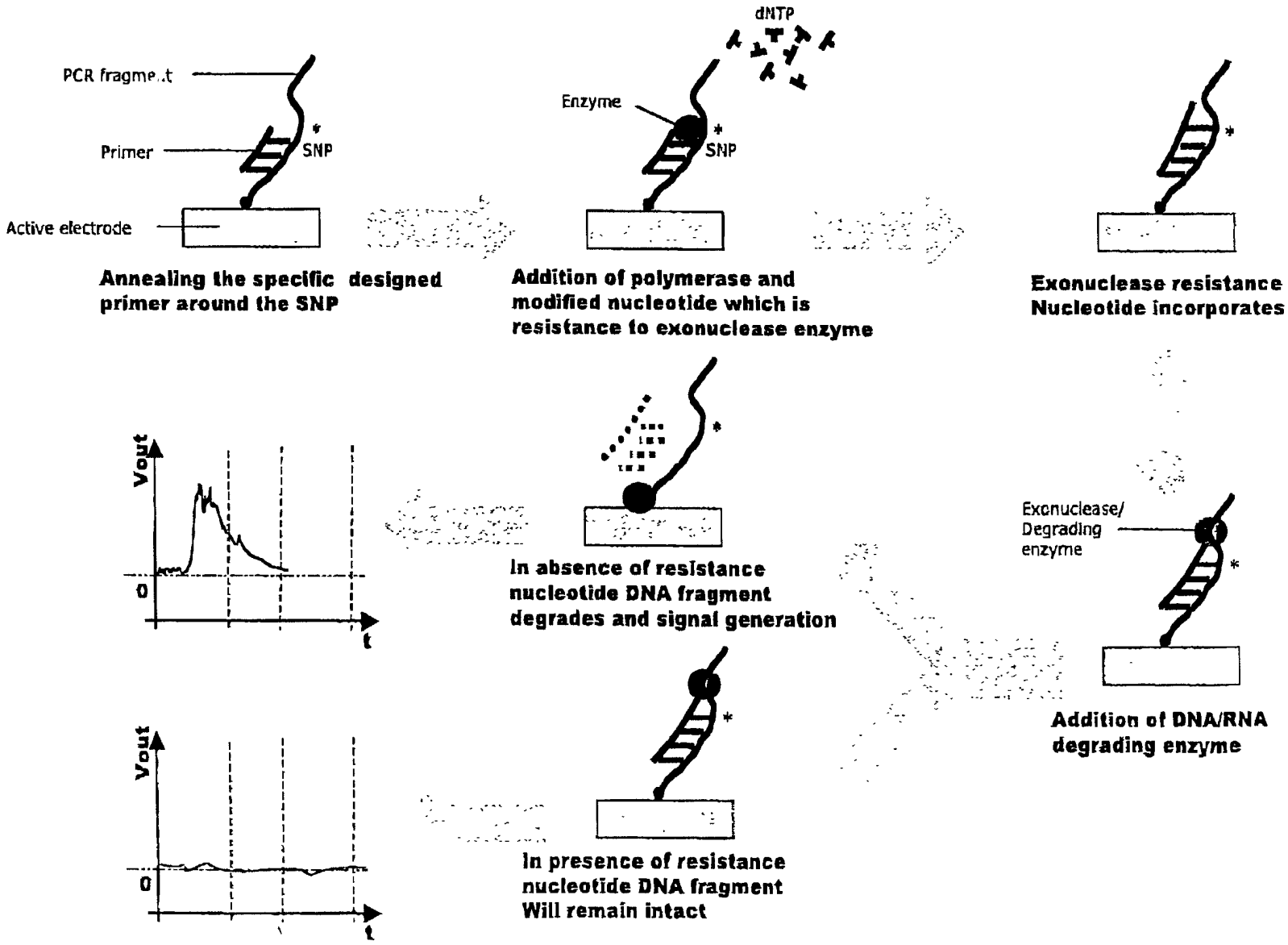


FIGURE 7:
Real time PCR

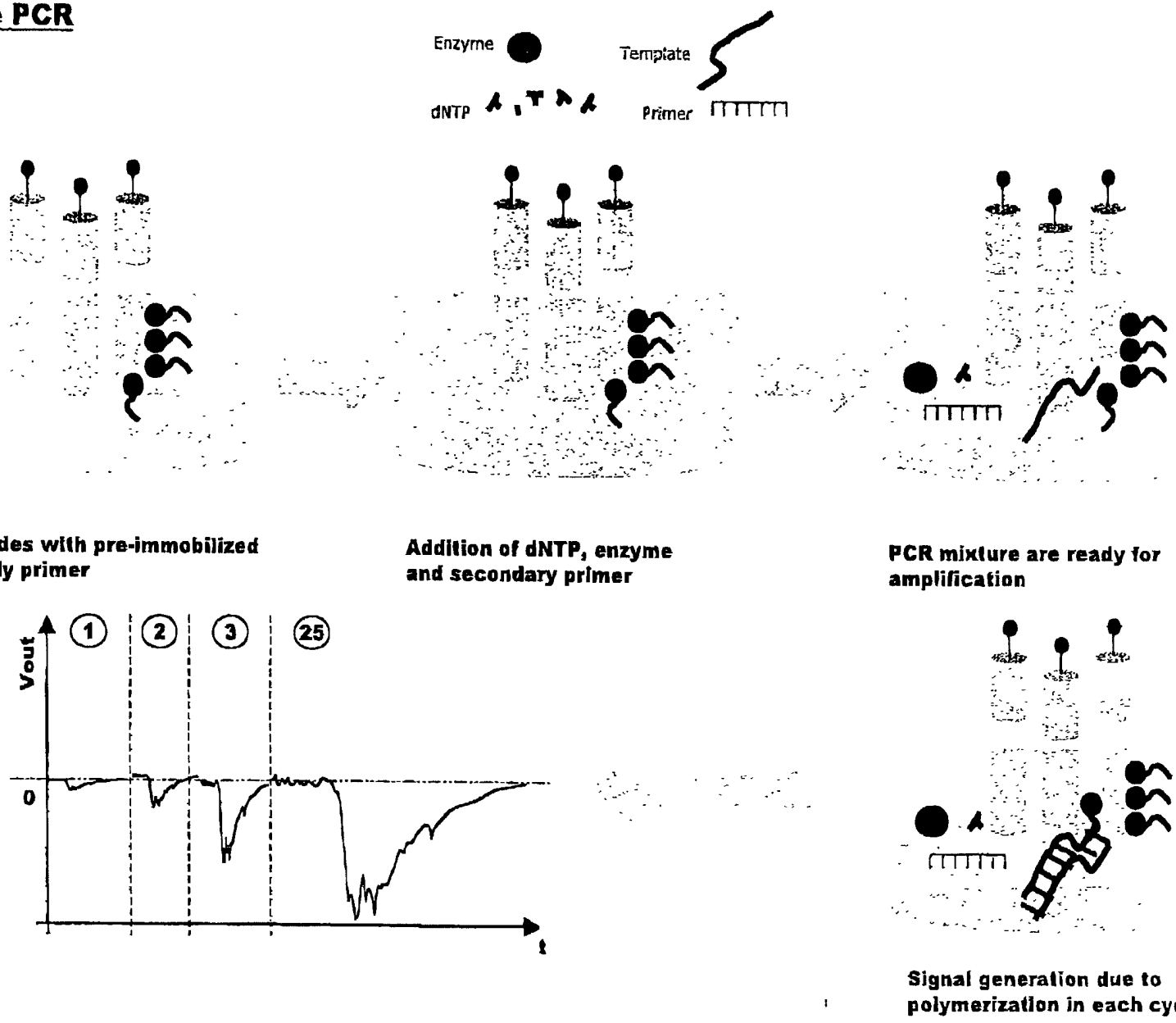
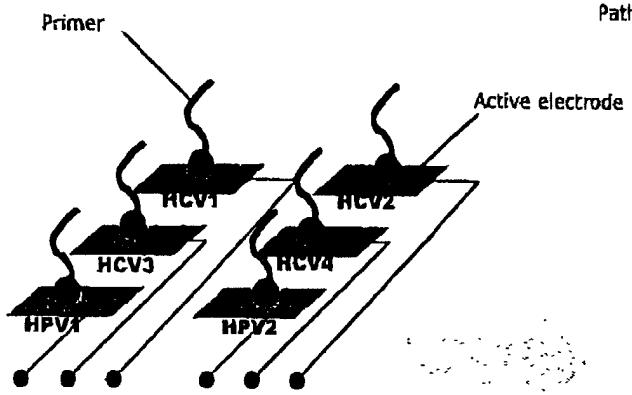
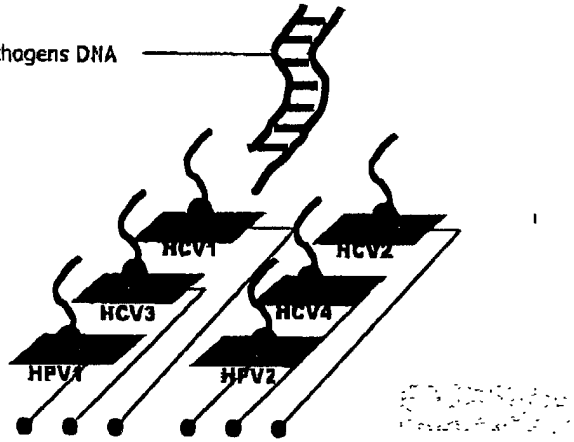


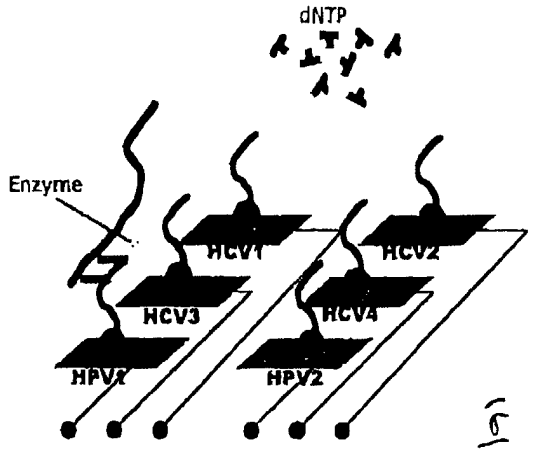
FIGURE 8:
Pathogen typing



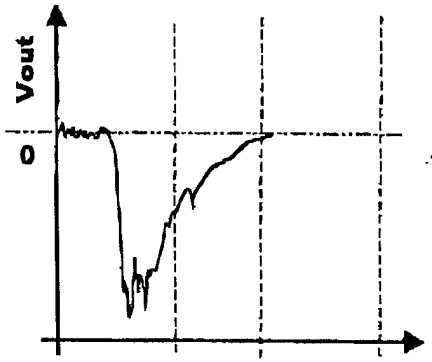
Immobilization of Pathogen specific primer

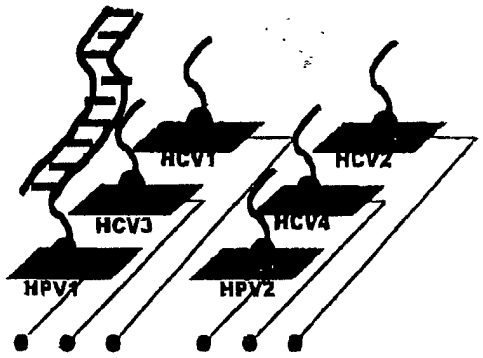


Denaturing and hybridization of pathogens DNA/RNA



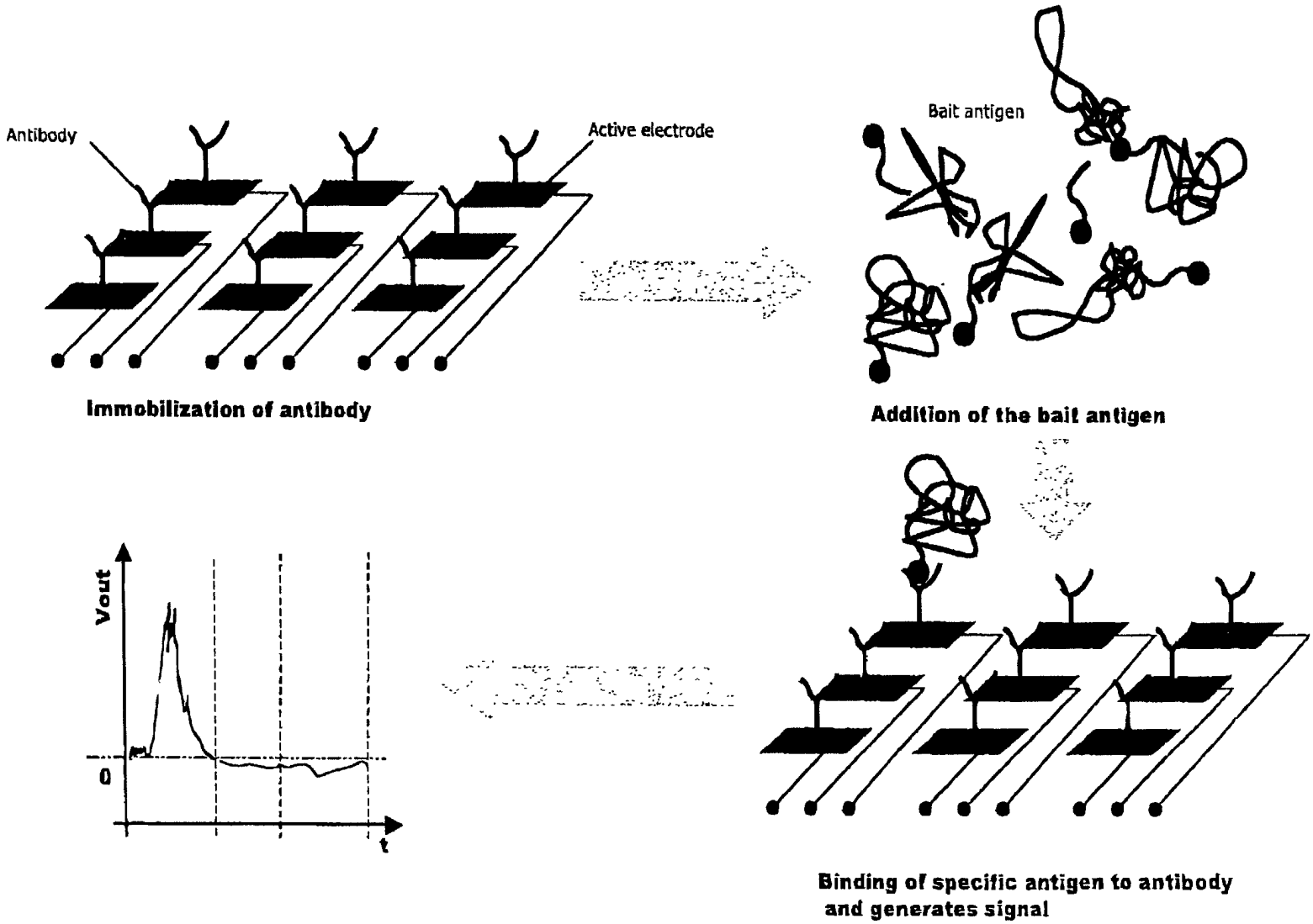
Addition of enzyme and dNTP





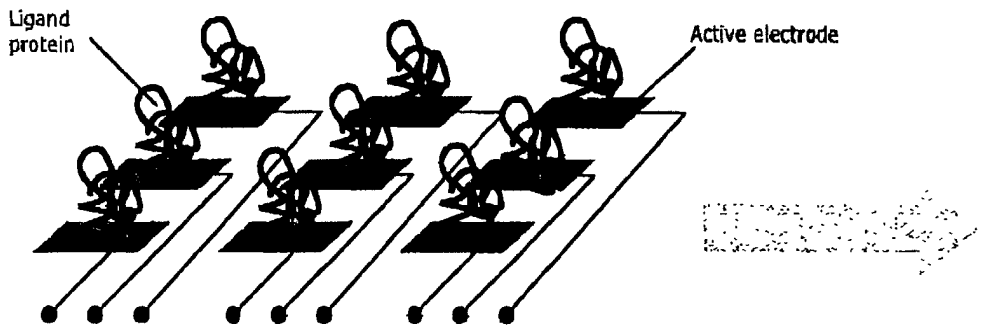
Only the hybridized primer will polymerize and generates signal

FIGURE 9:
Antigen-antibody detection

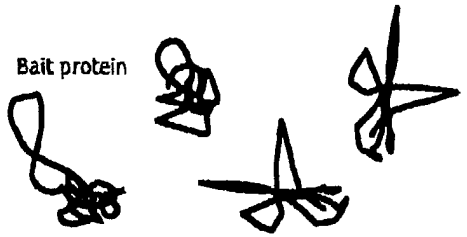


FEB-26-02 TUE 12:55 PM BOZICEVIC FIELD&FRANCIS FAX NO. 650 327+3231

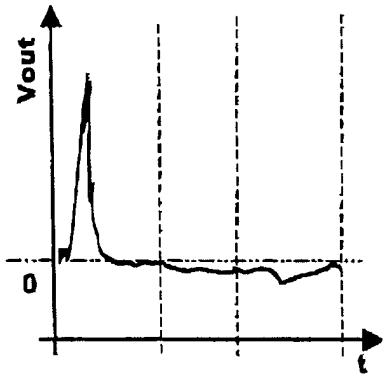
FIGURE 10:
Protein-protein interaction

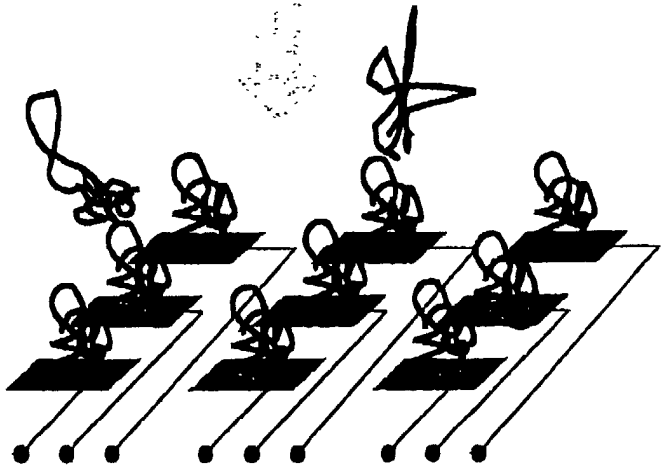


Immobilization of different ligand protein



Addition of the bait proteins





Specific protein-protein interaction and generates signal

FEB-26-02 TUE 12:56 PM BOZICEVIC FIELD&FRANCIS FAX NO. 650 327+3231

FIGURE 11:
Ligand and receptor detection

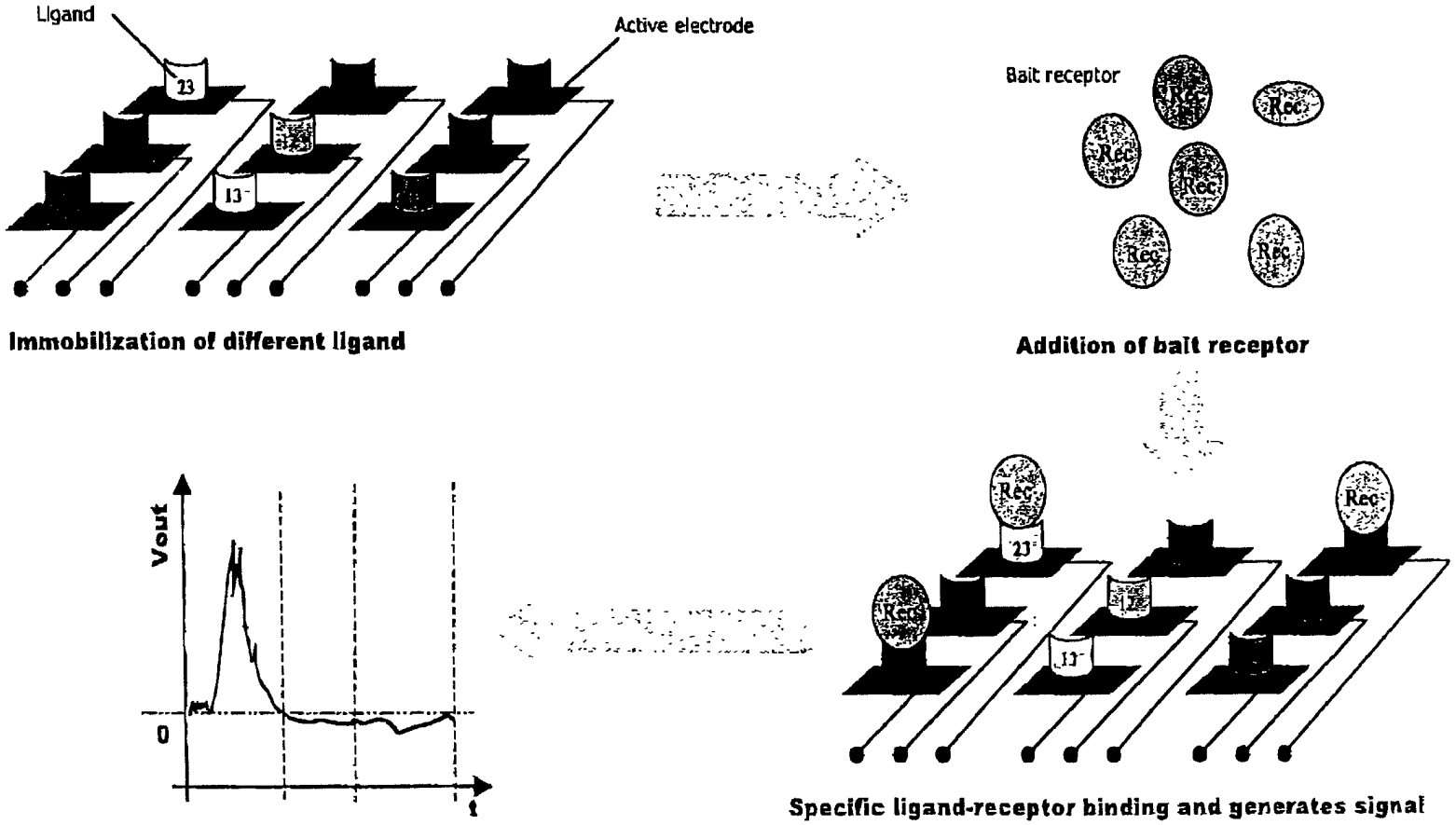
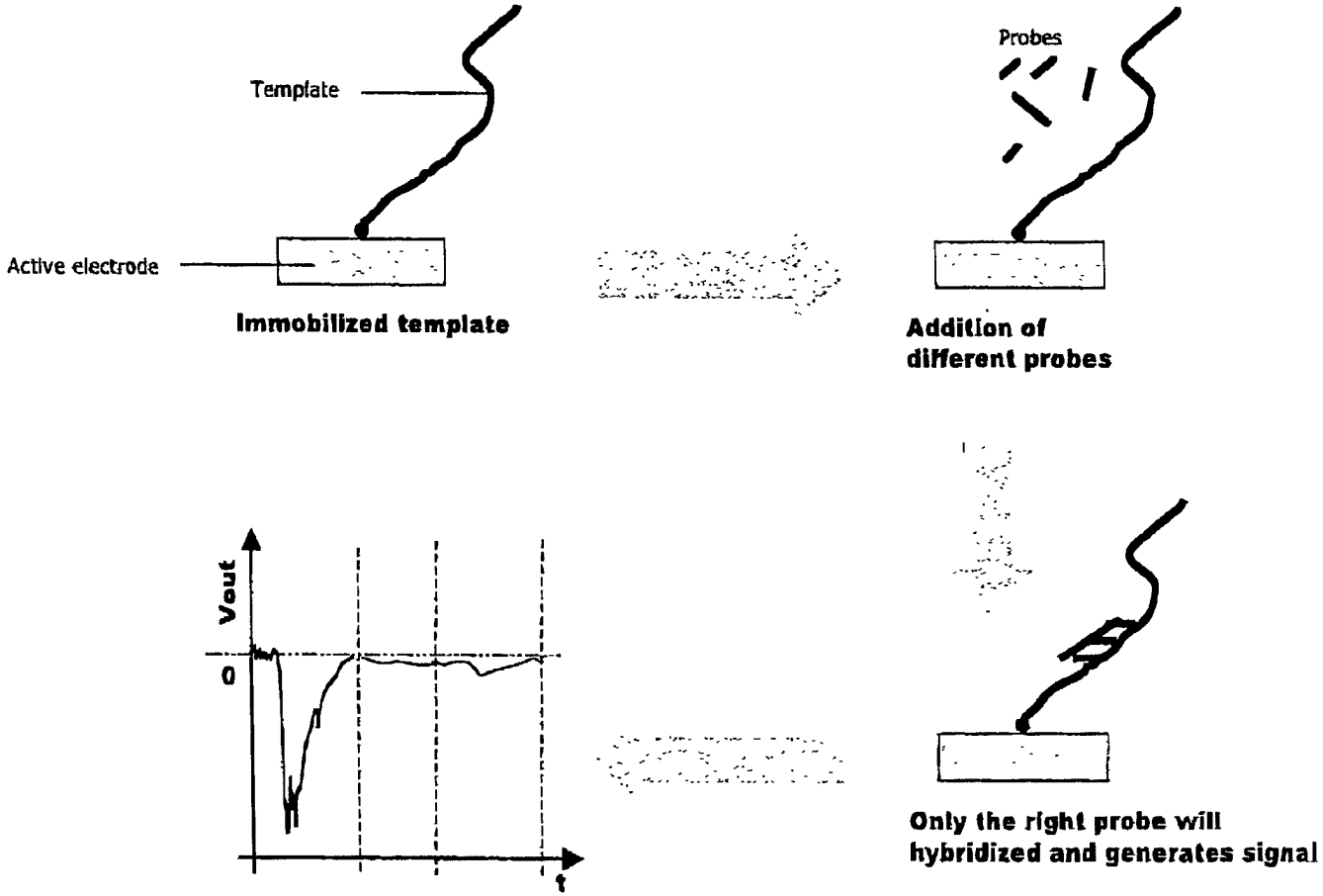


FIGURE 12:
Hybridization



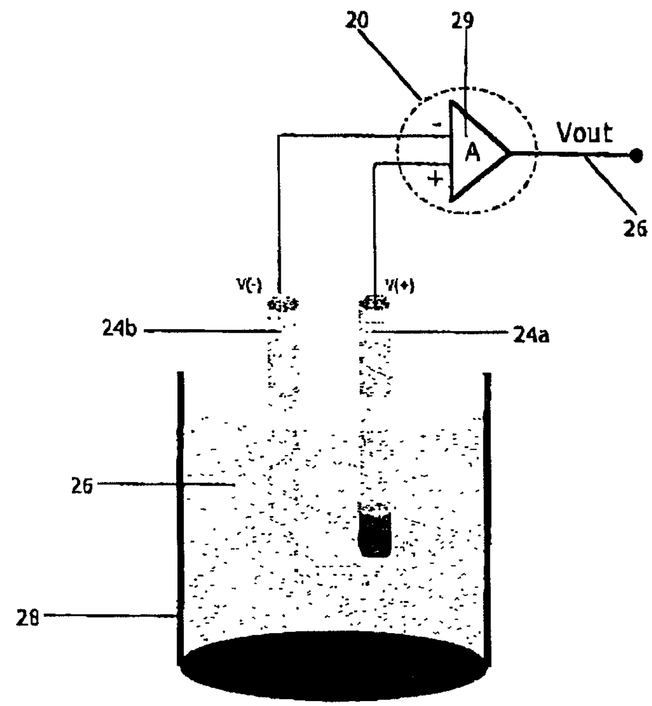


Fig: 13

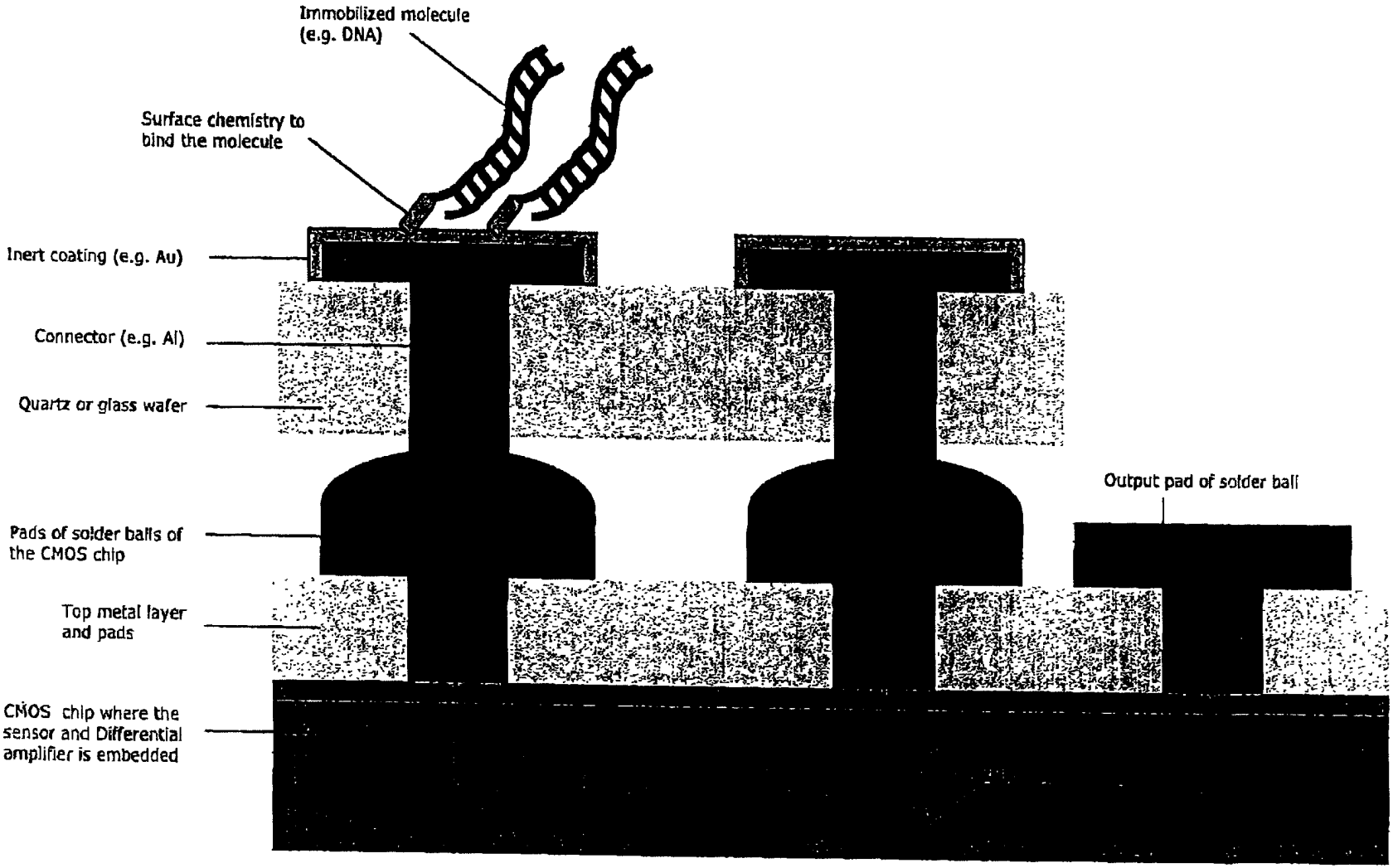


Fig: 14

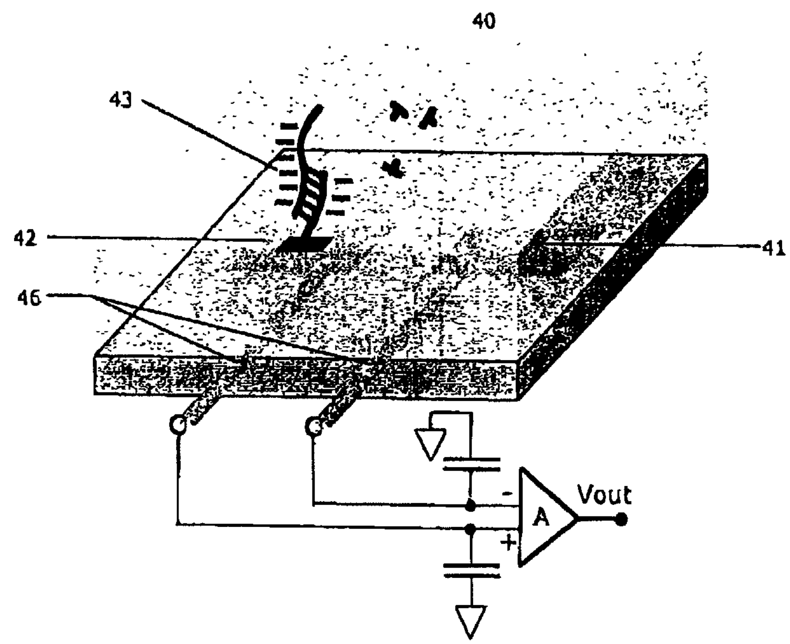


Fig: 15

FEB-26-02 TUE 12:57 PM BOZICEVIC FIELD&FRANCIS FAX NO. 650 327+3231 P. 25

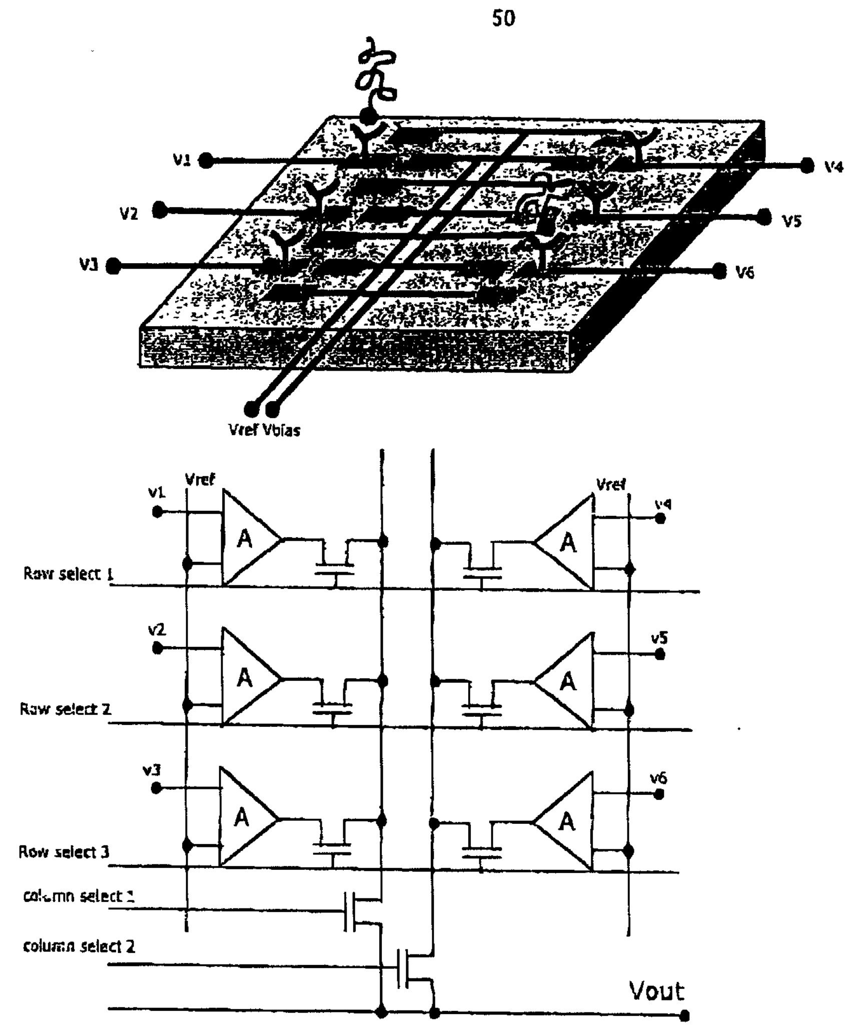
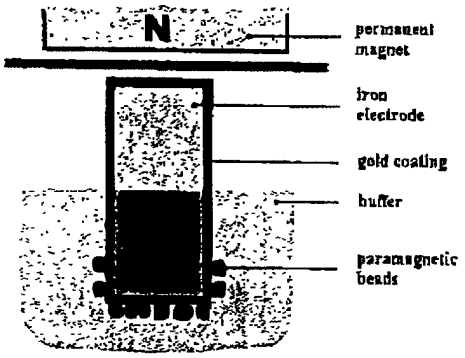


Fig: 16

Figure 17A: PCR product attracts to an electrode by using a permanent magnet and paramagnetic beads.



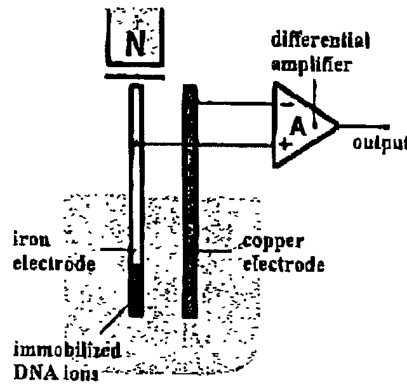


Figure 17 B: Basic model of the sensor with a differential amplifier

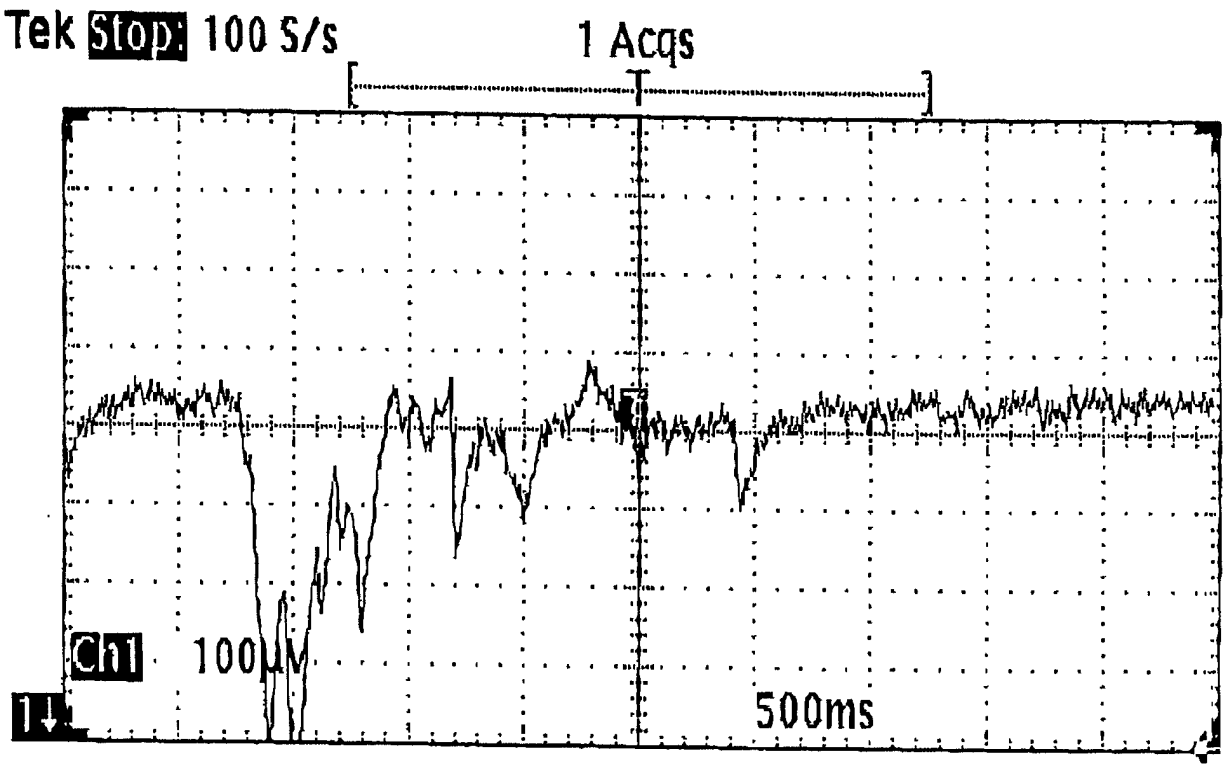


Figure 18 A: some sample charge sequencing extension signatures for 300 bp DNA

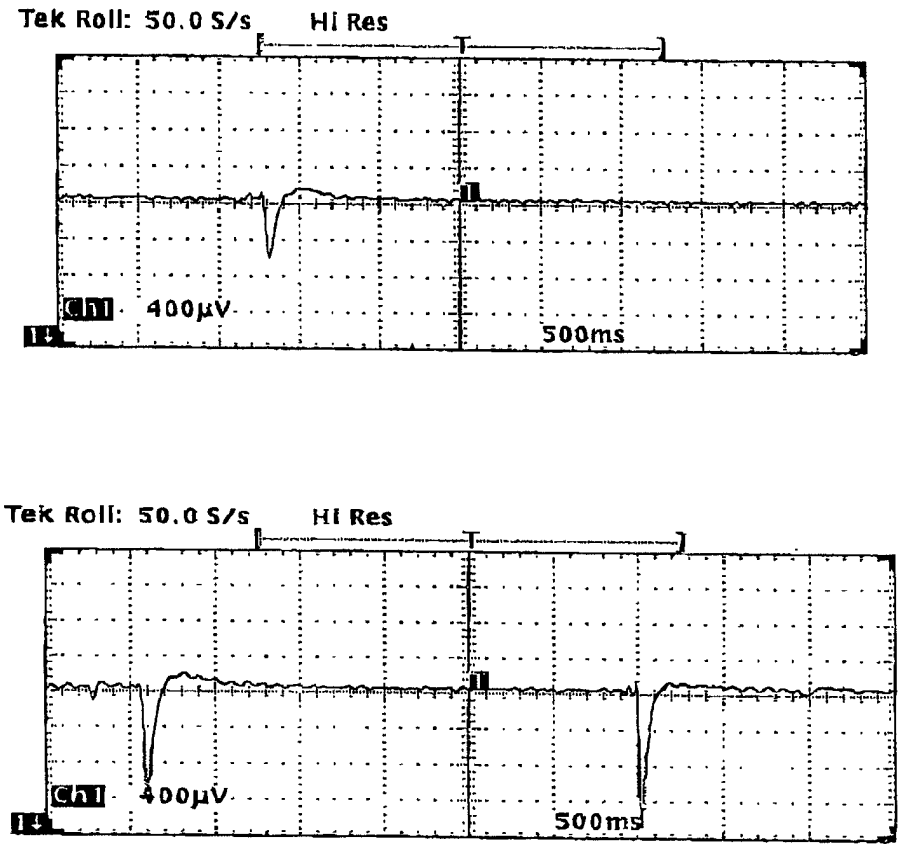


Figure 18 B: More sample charge sequencing extension signatures for 300 bp DNA with two different concentration of immobilized DNA (0.05 pmol and 0.1 pmol)

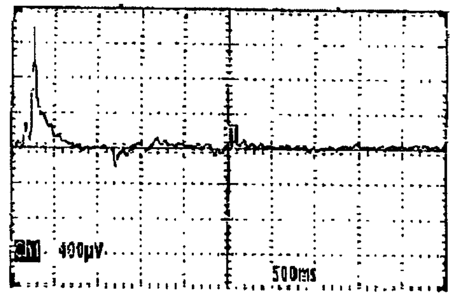


Figure 18C.

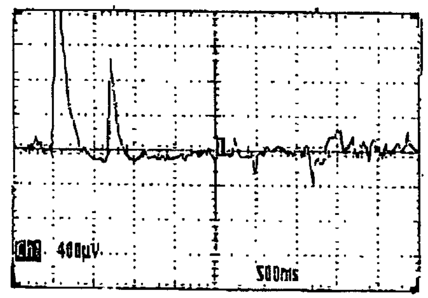


Figure 18D.

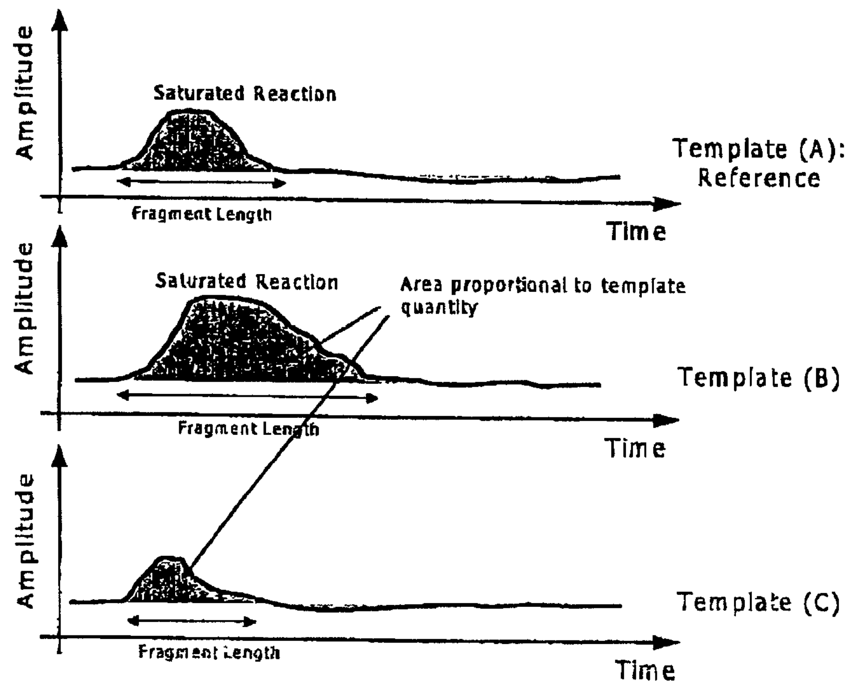
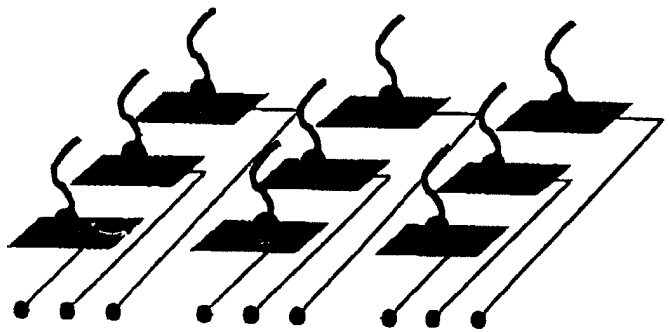

Amplitude
Saturated Reaction
Fragment Length
Template (A): Reference
Time
Amplitude
Saturated Reaction
Area proportional to template quantity
Fragment Length
Template (B)
Time
Amplitude
Fragment Length
Template (C)
Time

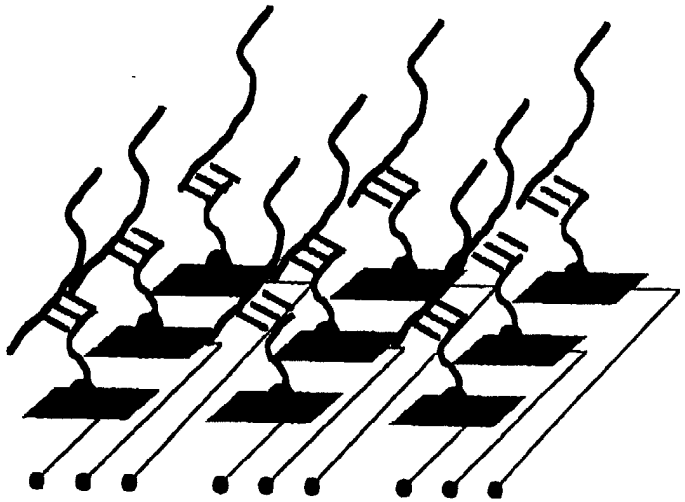
FIGURE 18E

FIGURE 19 a:

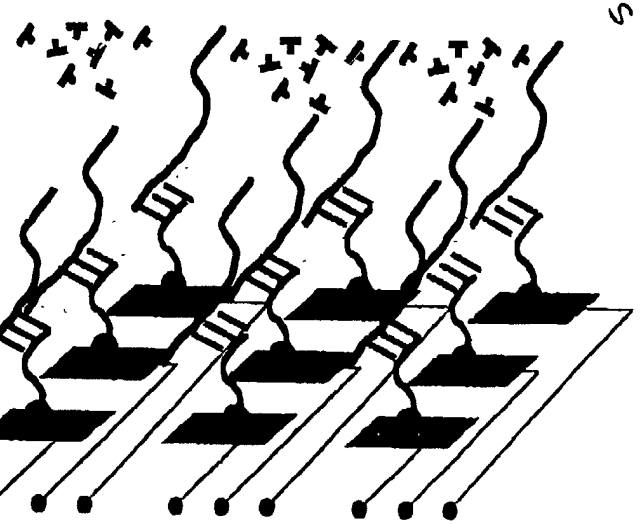
Gene Expression profiling by using Total RNA



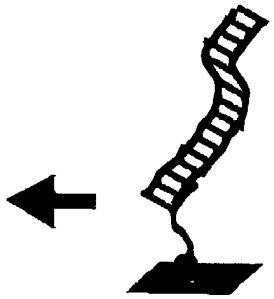
Gene specific designed primer



Hybridization of RNA to Specific primer



Addition of enzyme and nucleotide



Right nucleotide added and Generates signal

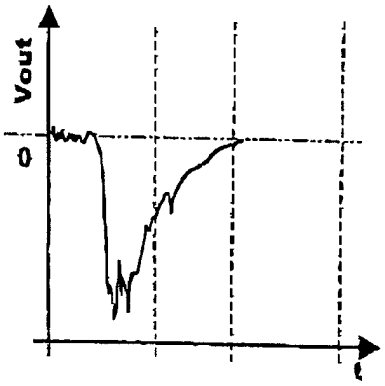
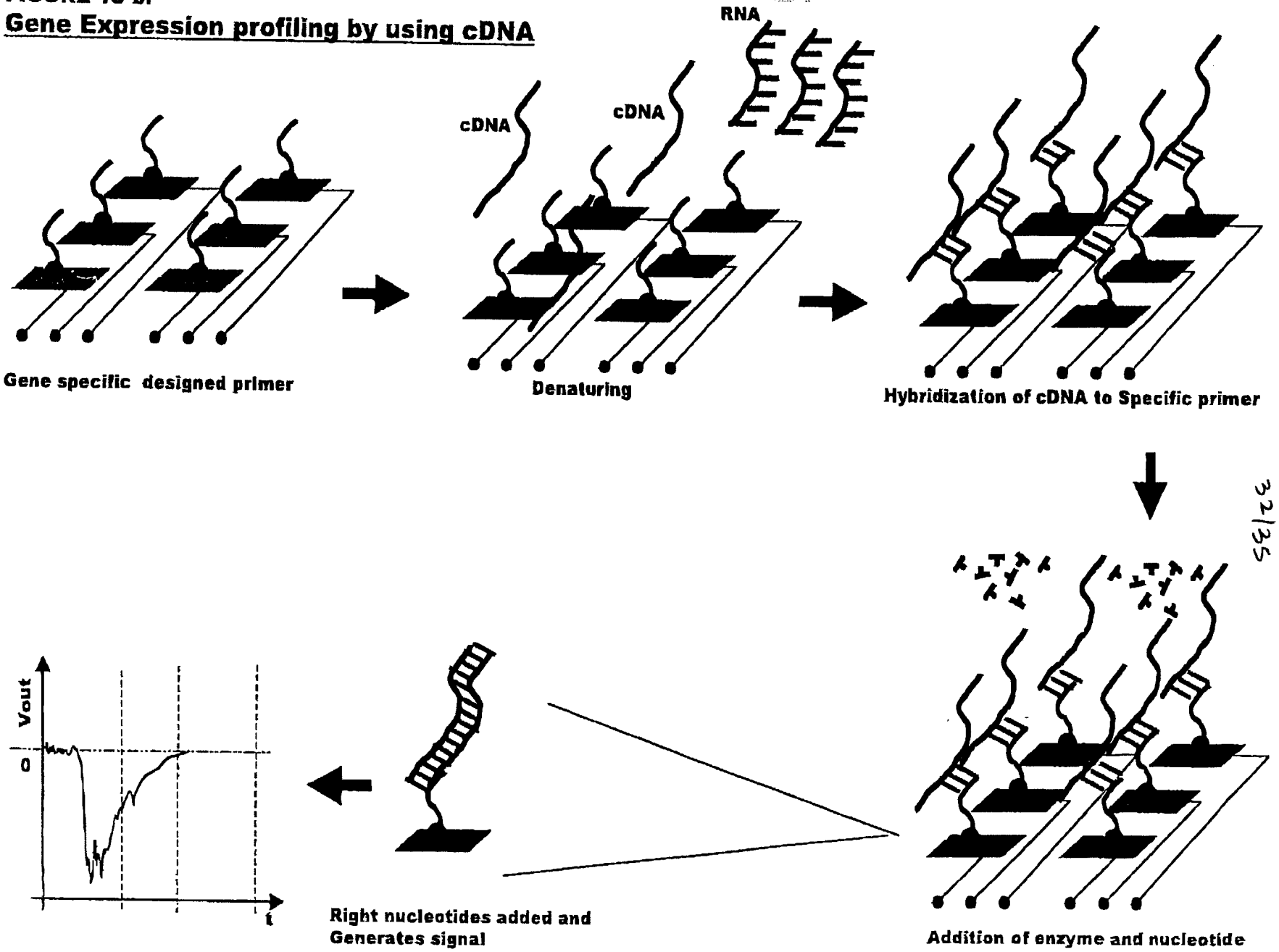


FIGURE 19 b:

Gene Expression profiling by using cDNA



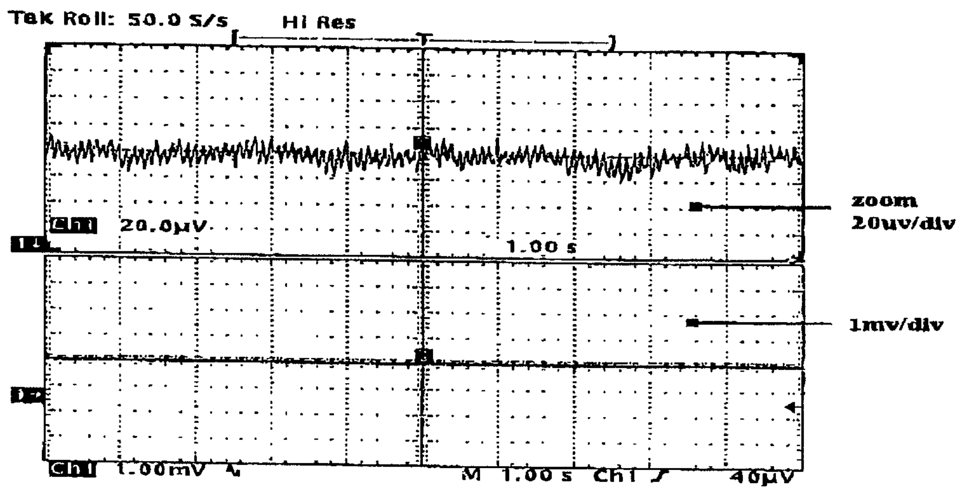

Tek Roll: 50.0 S/s
Hi Res
Ch1 20.0µV
1.00 s
zoom
20uv/div
1mv/div
Ch1 1.00mV
M 1.00 s Ch1
40µV

Figure 20

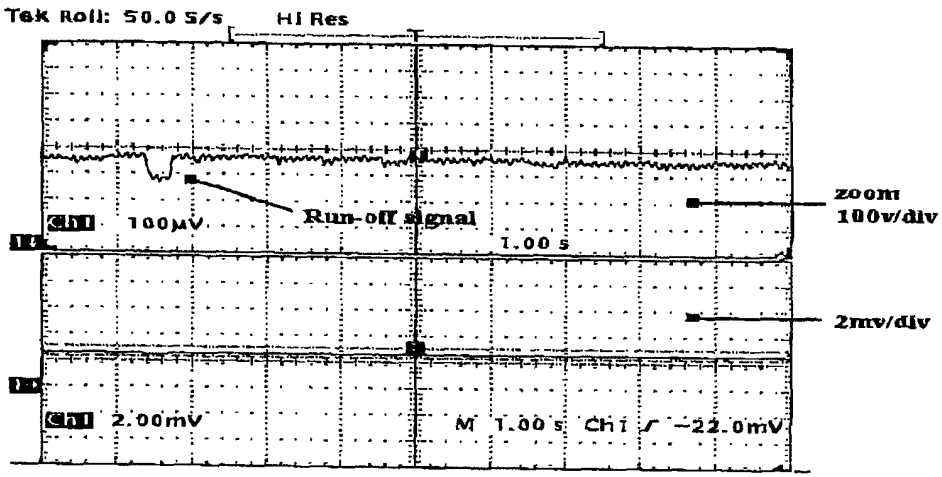
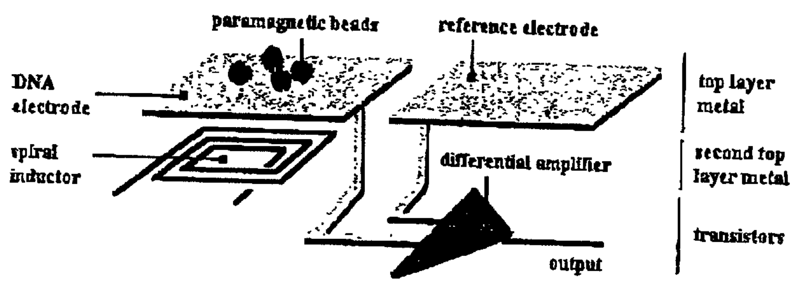

Tek Roll: 50.0 S/s
Hi Res
Run-off signal
zoom
100v/div
Ch1 100µV
1.00 s
2mv/div
Ch1 2.00mV
M 1.00 s Ch1 ƒ -22.0mV

Figure 21

Figure 22


paramagnetic beads
reference electrode
DNA electrode
top layer metal
spiral inductor
differential amplifier
second top layer metal
transistors
output